INFORMATION DISCLOSURE CITATION (Use several sheets if necessary) PTO Form 1449 December 4, 2008	Attorney Docket No. **056291-5283**	Application No. **10/578,663**
	Applicants: **HENNEQUIN *et al.***	
	Filing Date: **January 17, 2007**	Group Art Unit: **1624**

U.S. PATENT DOCUMENTS

Initial		Document No.	Date	Name	Class	Sub-Class	Filing Date
	1.	US 2003/0186995	October 2, 2003	Kath et al.			
	2.	US 2004/0048880	March 11, 2004	Himmelsbach et al.			

FOREIGN PATENT DOCUMENTS

		Document No.	Date	Country	Class	Sub-Class	Translation
	3.	CA 2476008	October 9, 2003	Canada			
	4.	CA 2543649	May 12, 2005	Canada			
	5.	WO 01/21596	March 29, 2001	WIPO			
	6.	WO 2004/046101	June 3, 2004	WIPO			
	7.	WO 2004/006846	January 22, 2004	WIPO			
	8.	WO 2005/013998	February 17, 2005	WIPO			
	9.	WO 2005/041973	May 12, 2005	WIPO			
	10.	WO 2005/097134	October 20, 2005	WIPO			

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	11.	Ballard et al. "Developing a small molecule erbB2 inhibitor:challenges with optimising DMPK properties" Poster - Presented at DMDG Cambridge (February 6, 2008)
	12.	Ballard et al. "Neutral 5-substituted 4-anilinoquinazolines as potent, orally active inhibitors of erbB2 receptor tyrosine kinase" Bioorg Med Chem Lett. 17(22):6326-6329 (2007)
	13.	Barlaam et al. "A new series of neutral 5-substituted 4-anilinoquinazolines as potent, orally active inhibitors of erbB2 receptor tyrosine kinase" Bioorganic & Medicinal Chemistry Letters 18(2):674-678 (2008)
	14.	Barlaam et al. "Indazolylamino/Anilinoquinazolines Bearing a C-5 substitution as erbB2 kinase inhibitors: Structure-activity relationships and identification of a candidate drug" at AACR in 2007
	15.	Barlaam et al. "Neutral 5-substituted 4-indazolylaminoquinazolines as potent, orally active inhibitors of erbB2 receptor tyrosine kinase" Bioorganic & Medicinal Chemistry Letters 18(6):1799-1803 (2008)
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	17.	Cockerill et al. "Indazolylamino quinazolines and pyridopyrimidines as inhibitors of the EGFr and c-erbB-2" Bioorganic & Medicinal Chemistry Letters 11(11):1401-1405 (2001)
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	19.	Gaul et al. "Discovery and Biological Evaluation of Potent Dual ErbB-2/EGFR Tyrosine Kinase Inhibitors: 6-Thiazolylquinazolines" Bioorganic & Medicinal Chemistry Letters 13(4):637-640 (2003)
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Examiner	Date Considered

Examiner: Initial if reference considered, whether or not citation is in conformance with MPEP 609; draw line through citation if not in conformance and not considered. Include copy of this form with next communication to applicant.

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	22.	Jani et al. "Discovery and pharmacologic characterization of CP-724,714, a selective ErbB2 tyrosine kinase inhibitor" Cancer Research 67(20):9887-9893 (2007)
	23.	Klutchko et al. "Tyrosine kinase inhibitors. 19. 6-Alkynamides of 4-anilinoquinazolines and 4-anilinopyrido[3,4-d]pyrimidines as irreversible inhibitors of the erbB family of tyrosine kinase receptors" J Med Chem. 49(4):1475-1485 (2006)
	24.	Petrov et al. "Optimization and SAR for dual ErbB-1/ErbB-2 tyrosine kinase inhibition in the 6-furanylquinazoline series" Bioorg Med Chem Lett. 16(17):4686-4691 (2006)

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